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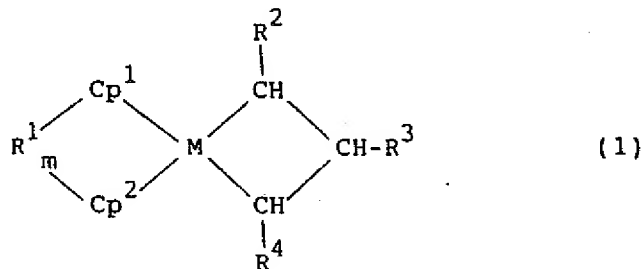
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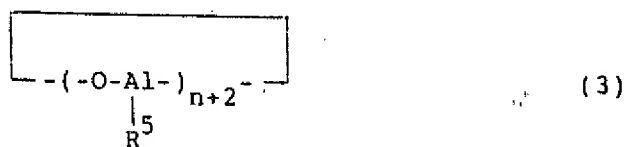
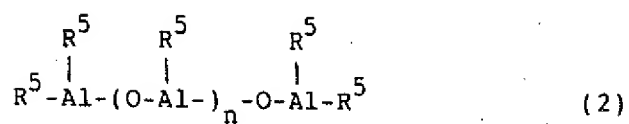
(54) Catalyst for the polymerization of vinyl compounds.

(57) A novel catalyst for polymerization of a vinyl compound is provided which comprises, as a catalyst component (A), an organometal complex represented by general formula (1) below:



where Cp<sup>1</sup> and Cp<sup>2</sup> are independently a substituted or unsubstituted cyclopentadienyl group; R<sup>1</sup> is an alkylene or arylalkylene group having 1 to 20 carbons, a dialkylsilylene, dialkylgermylene, alkylphosphinediyl, or alkylimino group, linking Cp<sup>1</sup> and Cp<sup>2</sup>; m is 0 or 1; M is titanium, zirconium, or hafnium; and R<sup>2</sup>, R<sup>3</sup>, and R<sup>4</sup> are independently hydrogen, a hydrocarbon group of 1 to 12 carbons, an alkoxy or aryloxy group, and a catalyst component (B) represented by the general formula (2) or (3)

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where  $n$  is an integer of from 4 to 60, and  $\text{R}^5$  is a hydrocarbon group. A process for producing a vinyl compound polymer which employs the catalyst defined above is also provided.

The present invention relates to a catalyst for producing a polymer of a vinyl compound (including  $\alpha$ -olefin polymers and aromatic polymers), and a process for producing a vinyl polymer employing the catalyst. More particularly, the present invention relates to a process for producing an aromatic vinyl polymer having mainly syndiotactic structure with high catalyst activity and with high selectivity.

Aromatic vinyl polymers include three structure types of polymers: syndiotactic polymers, isotactic polymers, and atactic polymers.

Of these, the polymers of the syndiotactic structure, which have a high melting point and crystallize quickly in comparison with polymers of other structures, are useful as heat-resistant polymers. The syndiotactic aromatic vinyl polymer is produced, for example, in the presence of a catalyst formed by contact of a titanium compound such as titanium halide and alkoxytitanium with an organoaluminum compound such as methylalumoxane as disclosed in JP-A- 62-04818 (1987).

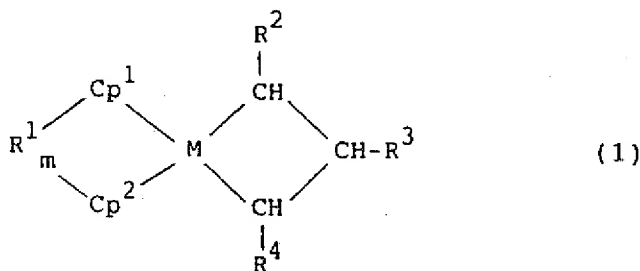
However, in the polymerization of styrene with a catalyst system comprising the combination of a titanium compound such as titanium tetrachloride and tetraethoxytitanium with methylalumoxane, the catalyst activity is low, and the catalyst remains in the formed polymer in a considerable amount. Therefore the polymer is presumed to discolor significantly during high-temperature molding, and not to be suitable for practical use.

On the other hand, a catalyst system composed of methylaluminoxane and a complex obtained by reaction of a transition metal compound, like titanium tetrachloride, with an organic compound, like 2,2'-dihydroxy-3,3'-ditert-butyl-5,5'-dimethyl-diphenylsulfide, gives a slightly lower content of stereoregular polymers owing mainly to atactic polymer formation as a by-product even though the catalyst exhibits considerably high catalytic activity. The amorphous polymer coexisting in a larger amount affects adversely the melting point and the crystallization velocity of the polymer. Therefore, as a disadvantage the removal of the amorphous polymer is required by solvent extraction or the like.

Thus, it is the object of the present invention to provide a catalyst which enables the production of aromatic vinyl polymers of syndiotactic structure with high catalyst activity and high selectivity.

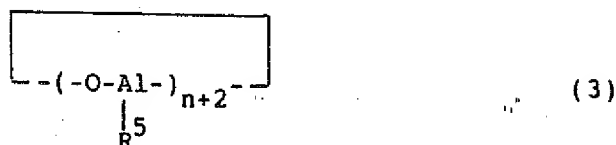
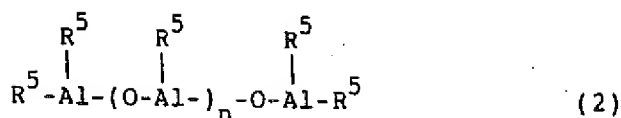
This object has been achieved by a specific organometal complex in combination with methylaluminoxane.

The catalyst for polymerization of a vinyl compound of the present invention comprises, as a catalyst component, a novel organometal complex represented by the general formula (1) below:



where  $\text{Cp}^1$  and  $\text{Cp}^2$  are independently a substituted or unsubstituted cyclopentadienyl group;  $\text{R}^1$  is an alkylene or arylalkylene group having 1 to 20 carbons, a dialkylsilylene, dialkylgermanylene, alkylphosphinediyl, or alkylimino group, linking  $\text{Cp}^1$  and  $\text{Cp}^2$ ;  $m$  is 0 or 1;  $\text{M}$  is titanium, zirconium, or hafnium; and  $\text{R}^2$ ,  $\text{R}^3$ , and  $\text{R}^4$  are independently hydrogen, a hydrocarbon group of 1 to 12 carbons, an alkoxy or aryloxy group.

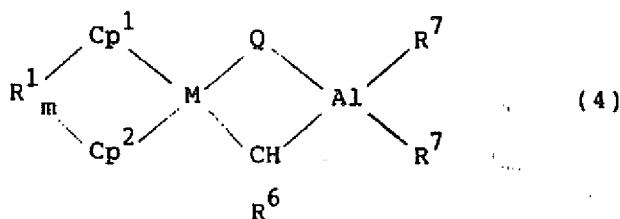
The catalyst for the polymerization of a vinyl compound of the present invention comprises a catalyst component (A) represented by the general formula (1) above, and a catalyst component (B) represented by the general formula (2) or (3)



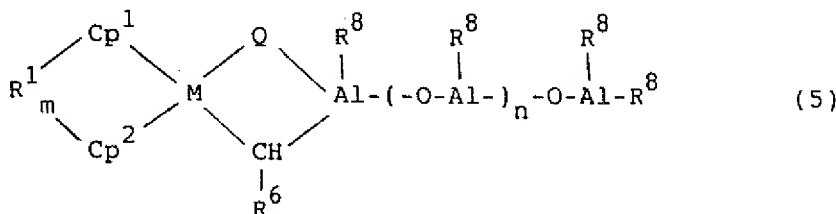
where  $n$  is an integer of from 4 to 60, and  $\text{R}^5$  is a hydrocarbon group.

The present invention further provides a process for producing stereoregular aromatic vinyl polymer of high syndiotacticity with high selectivity by use of the above catalyst.

The catalyst component (A) of the catalyst for polymerization of a vinyl compound of the present invention can be prepared by reacting the organometal compound represented by the general formula (4) or (5) with an  $\alpha$ -olefin represented by the general formula (6). General Formula (4):



where  $\text{Cp}^1$  and  $\text{Cp}^2$  are independently a substituted or unsubstituted cyclopentadienyl group;  $\text{R}^1$  is an alkylene or arylalkylene group having 1 to 20 carbons, a dialkylsilylene, dialkylgermanylenes, alkylphosphinediyl, or alkylimino group, linking  $\text{Cp}^1$  and  $\text{Cp}^2$ ;  $m$  is 0 or 1;  $\text{Q}$  is a hydrocarbon group of 1 to 12 carbons or a halogen atom;  $\text{R}^6$  is hydrogen, a hydrocarbon group of 1 to 12 carbons, an alkoxy or aryloxy group;  $\text{R}^7$  is hydrogen or a hydrocarbon group of 1 to 12 carbons, and  $\text{M}$  is titanium, zirconium, or hafnium. General Formula (5):



where  $\text{Cp}^1$  and  $\text{Cp}^2$  are independently a substituted or unsubstituted cyclopentadienyl group;  $\text{R}^1$  is an alkylene or arylalkylene group having 1 to 20 carbons, a dialkylsilylene, dialkylgermanylenes, alkylphosphinediyl, or alkylimino group, linking  $\text{Cp}^1$  and  $\text{Cp}^2$ ;  $m$  is 0 or 1;  $\text{Q}$  is a hydrocarbon group of 1 to 12 carbons or a halogen atom;  $\text{R}^6$  is hydrogen, a hydrocarbon group of 1 to 12 carbons, an alkoxy or aryloxy group;  $\text{R}^8$  is hydrogen or a hydrocarbon group of 1 to 12 carbons;  $n$  is an integer of from 4 to 60; and  $\text{M}$  is titanium, zirconium, or hafnium. General Formula (6):



5 where R<sup>9</sup> is hydrogen or a hydrocarbon group of 1 to 12 carbons.

The component represented by the general formula (4) used for synthesis of the catalyst component (A) of the present invention includes specifically (μ-chloro)(μ-methylene)bis(cyclopentadienyl)-(dimethylaluminum)titanium, (μ-chloro)(μ-methylene)methylenebis(cyclopentadienyl)(dimethylaluminum)-  
10 titanium, (μ-chloro)(μ-methylene)dimethylsilylbis(cyclopentadienyl)(dimethylaluminum)titanium, and (μ-chloro)(μ-methylene)isopropylidenebis(cyclopentadienyl)(dimethylaluminum)titanium.

The component represented by the general formula (5) used for synthesis of the catalyst component (A) of the present invention includes bis(cyclopentadienyl)titanium-methylaluminoxane complex, methylenebis(cyclopentadienyl)titanium-methylaluminoxane complex, dimethylsilylbis(cyclopentadienyl)-  
15 titanium-methylaluminoxane complex, and isopropylidenebis(cyclopentadienyl)titanium-methylaluminoxane complex.

The component represented by the general formula (6) used for synthesis of the catalyst component (A) of the present invention includes e.g. ethylene, propylene, 1-butene, 1-hexene, 1-octene, styrene, methystyrene, chlorostyrene and methoxystyrene.

20 The reaction of the compound of the general formula (4) or (5) with the compound of the general formula (6) is conducted generally in the presence of a solvent.

The molar ratio of the compound of the general formula (4) or (5) to the compound of the general formula (6) is not limited. However, the molar ratio of the compound of the general formula (4) to the compound of the general formula (6) is preferably in the range of from 1:0.5 to 1:10, more preferably from  
25 1:1 to 1:3. The molar ratio of the compound of the general formula (5) to the compound of the general formula (6) is preferably in the range of from 1:0.5 to 1:30, more preferably from 1:1 to 1:10.

The solvent used includes halogenated hydrocarbons such as chloroform and carbon tetrachloride, and aromatic hydrocarbons such as benzene, toluene, and xylene.

30 The reaction temperature depends on the starting material, the solvent, and other conditions, and is usually in the range of from -50 to 60 °C.

The intended compound can be isolated in high purity from the resulting reaction mixture by removing the solvent by vacuum evaporation and recrystallizing the evaporation residue from an organic solvent such as ether.

35 The catalyst component (A) is confirmed to have the structure of the general formula (1) by proton nucleomagnetic resonance spectroscopy.

The catalyst component (B) is an aluminoxane represented by the general formula (2) or (3). The substituent on the aluminum of the aluminoxane is a hydrocarbon group of 1 to 6 carbons such as methyl, ethyl, propyl, and butyl; preferably methyl. The oligomerization degree is from 6 to 62. This type of compound may be prepared by a known method, for example, by causing reaction by adding an aluminum  
40 compound into a suspension of a hydrous salt (e.g. copper sulfate hydrate, aluminum sulfate hydrate, etc.) in a hydrocarbon medium.

The molar ratio of the catalyst component (B) to the catalyst component (A), namely (B)/(A), is in the range of from 10 to 1000.

45 The vinyl compound polymerizable according to the present invention includes α-olefins, styrene, and derivatives thereof. The derivatives of styrene include alkylstyrenes such as methylstyrene, ethylstyrene, and dimethylstyrene; halogenated styrenes such as chlorostyrene, bromostyrene, and fluorostyrene; halogen-substituted alkylstyrenes such as chloromethylstyrene; alkoxy styrenes such as methoxystyrene; carboxymethylstyrene, and alkylsilylstyrene.

50 The vinyl compound is polymerized in the presence of the above catalyst. The polymerization may be conducted in bulk, or in an aliphatic hydrocarbon such as pentane, hexane, or heptane, or in an aromatic hydrocarbon such as benzene, toluene, and xylene.

The concentration of the catalyst component used in the solution is preferably in the range of from 0.1 to 1000 mmol/l. The polymerization temperature is not specially limited, but is usually in the range of from -78 to 150 °C.

55 The present invention is described in more detail by reference to the examples.

Example 1

## Synthesis of Methylenebis(cyclopentadienyl)-2-phenyltitanacyclobutane Complex:

- 5 One gram of ( $\mu$ -chloro)( $\mu$ -methylene)methylenebis(cyclopentadienyl)(dimethylaluminum)titanium was dissolved in 6 ml of toluene, and thereto 0.36 g of styrene was added. The mixture was stirred at room temperature. Then 0.47 g of dimethylaminopyridine was added to the reaction system, whereby a precipitate was formed gradually. The suspension was filtered with celite® to obtain a red solution. The solution was evaporated to dryness to obtain a reddish brown solid. This solid was dissolved in ether, and  
 10 left standing at -30 °C for 4 days. Thereby red needle-crystalline methylenebis(cyclopentadienyl)-2-phenyltitanacyclobutane was obtained in a yield of 0.3 g.

The resulting complex was identified by <sup>1</sup>H-NMR as follows: 0.1 ppm (m, -CH<sub>2</sub>-), 1.8 ppm (m, Ti-CH<sub>2</sub>-), 2.1 ppm (t, -(C<sub>6</sub>H<sub>5</sub>)CH-), 2.5 ppm (s, Cp-CH<sub>2</sub>-Cp), 4.7 ppm (t, Cp), and 6.8 ppm (t, Cp).

15 Example 2

## Synthesis of Methylenebis(cyclopentadienyl)-3-methyltitanacyclobutane Complex:

- The synthesis was conducted in the same manner as in Example 1 except that propylene was bubbled  
 20 into the solution in place the addition of styrene. Consequently, red crystalline methylenebis-(cyclopentadienyl)-3-methyltitanacyclobutane was obtained in a yield of 40 %.

The resulting complex was identified by <sup>1</sup>H-NMR as follows: 0.03 ppm (m, CH), 0.7 ppm (s, CH<sub>3</sub>), 2.3 - 3.2 ppm (m, Ti-CH<sub>2</sub>-C), 2.4 ppm (s, Cp-CH<sub>2</sub>-Cp), 4.7 ppm (t, Cp), and 6.7 ppm (t, Cp).

25 Example 3

## Synthesis of Methylenebis(cyclopentadienyl)-2-phenyltitanacyclobutane Complex:

- In 20 ml of toluene, 5 g of methylenebis(cyclopentadienyl)-methylaluminoxane complex was dissolved.  
 30 Styrene (10 equivalents) was added thereto, and the mixture was cooled to -20 °C with stirring. To the resulting red solution, a solution of methylaluminoxane (50 equivalent) in toluene was added dropwise gradually. The mixture was then brought to room temperature in 12 hours. The reaction solution was cooled to 0 °C. The insoluble matter was removed by filtration with celite®, and the filtrate was evaporated to dryness. The evaporation residue was dissolved in ether, and the solution was left standing at -40 °C for 5  
 35 days. As the result, red crystalline methylenebis(cyclopentadienyl)-2-phenyltitanacyclobutane in a yield of 0.8 g was isolated.

Example 4

- 40 In a nitrogen-purged Schlenk reaction vessel, 0.039 mmol of methylenebis(cyclopentadienyl)-2-phenyltitanacyclobutane prepared in Example 1 was placed. Thereto 10 ml of toluene was added. Further thereto, 6 ml of styrene was added. To the mixture, a solution of methylaluminoxane (16-mer) in toluene was added dropwise in an amount to give an Al/Ti molar ratio of 200. The reaction was allowed to proceed at 30 °C for 10 hours. Then 10 ml of methanol-hydrochloric acid solution was added to stop the reaction. The resulting  
 45 white polymer was collected by filtration, and dried to obtain 4.5 g of a polymer.

This polymer was extracted with methyl ethyl ketone by Soxhlet extraction. The polymer was found to contain a methyl ethyl ketone-soluble portion in an amount of 3 %.

- The melting point of the resulting polymer had a melting point of 267 °C by DSC measurement. The polymer had pentad rrrr of 97 %, according to <sup>13</sup>C-NMR structure analysis in o-dichlorobenzene, from the  
 50 peak of 145.5 ppm resulting from syndiotactic structure.

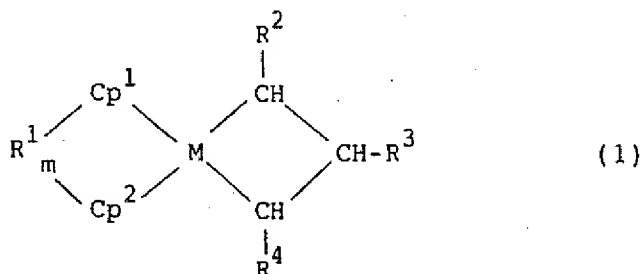
Comparative Example 1

- The polymerization was conducted in the same manner as in Example 4 except that 0.041 mmol of  
 55 methylenebis(cyclopentadienyl)titanium dichloride was used in place of methylenebis(cyclopentadienyl)-2-phenyltitanacyclobutane. As the result, the amount of the formed dry polymer was 0.9 g. This polymer was extracted with methyl ethyl ketone by Soxhlet extraction and was found to have a methyl ethyl ketone-soluble portion in an amount of 8 %

As shown above, the catalyst of the present invention enables the production of a highly syndiotactic aromatic vinyl compound polymer with high catalyst activity with high selectivity.

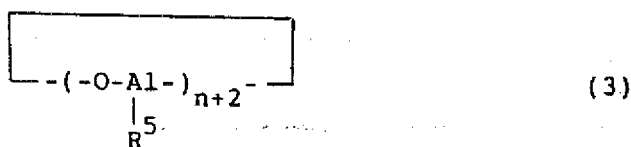
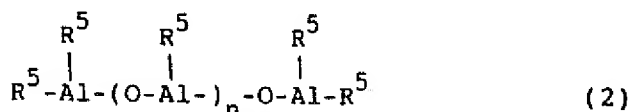
# Claims

1. An organometal complex represented by general formula (1) below:



where  $\text{Cp}^1$  and  $\text{Cp}^2$  are independently a substituted or unsubstituted cyclopentadienyl group;  $\text{R}^1$  is an alkylene or arylalkylene group having 1 to 20 carbons, a dialkylsilylene, dialkylgermanylenes, alkylphosphinediyl, or alkylimino group, linking  $\text{Cp}^1$  and  $\text{Cp}^2$ ;  $m$  is 0 or 1;  $\text{M}$  is titanium, zirconium, or hafnium; and  $\text{R}^2$ ,  $\text{R}^3$ , and  $\text{R}^4$  are independently hydrogen, a hydrocarbon group of 1 to 12 carbons, an alkoxy or aryloxy group.

2. A organometal complex according to claim 1, wherein  $\text{M}$  is titanium,  $\text{R}^2$  is phenyl, and  $\text{R}^3$  and  $\text{R}^4$  are each hydrogen.
3. A organometal complex according to claim 1, wherein  $\text{M}$  is titanium,  $\text{R}^3$  is methyl, and  $\text{R}^2$  and  $\text{R}^4$  are each hydrogen.
4. A catalyst for the polymerisation of a vinyl compound, comprising a catalyst component (A) of the general formula (1) as defined in any of claims 1 to 3, and a catalyst component (B) represented by the general formula (2) or (3)



where  $n$  is an integer of from 4 to 60, and  $\text{R}^5$  is a hydrocarbon group.

5. A process for producing a vinyl compound polymer which employs the catalyst defined in any of claims 1 to 4.



European Patent  
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# EUROPEAN SEARCH REPORT

Application Number  
EP 93 11 4423

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 5)
X	US-A-4 607 112 (ROBERT H. GRUBBS) * the whole document *	1	C07F17/00
X	J. AM. CHEM. SOC. vol. 102, 1980 pages 6876 - 6878 T.R. HOWARD 'Titanium Metallacarbene-Metallacyclobutane Reactions: Stepwise Metathesis'	1	
A	DE-A-11 91 374 (THE DOW CHEMICAL COMPANY) * the whole document *		
			TECHNICAL FIELDS SEARCHED (Int. Cl. 5)
			C07F C08F
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 15 December 1993	Examiner Fischer, B
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		I : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons A : member of the same patent family, corresponding document	

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